## Amendments to the Claims:

This listing of claims will replace all prior versions:

Claims 1-18 (canceled)

Claim 19 (previously presented): A molecule of general formula (I), and the pharmaceutically acceptable salts thereof:

$$(X_0)_{x0}$$
- $(X_1)_{x1}$ - $(X_2)_{x2}$ - $X_3$ - $(X_4)_{x4}$ - $X_5$ - $X_6$ - $(X_7)_{x7}$ - $(X_8)_{x8}$ - $(X_9)_{x9}$ 
(I)

in which

- x0, x1, x2, x4, x7, x8 and x9 each represent, independently, an integer equal to 0 or to 1;
- -X<sub>0</sub> represents a group:

with p ranging from 3 to 23;

- -X<sub>1</sub> and X<sub>3</sub> each represent a natural or synthetic amino acid in the L or D configuration, each comprising at least one hydroxyl function on its side chain;
- -X<sub>2</sub> represents a natural or synthetic amino acid in the L or D configuration chosen from those comprising an alkyl side chain;
- -X<sub>4</sub> represents a natural or synthetic amino acid in the L or D configuration chosen from those comprising an aromatic side chain;
- -X<sub>5</sub> represents an amino acid in the L or D configuration chosen from lysine, arginine, histidine, aspartic acid, asparagine, glutamic acid and glutamine:
- -X<sub>6</sub> represents an amino acid in the L or D configuration chosen from tyrosine, phenylalanine, leucine, isoleucine, alanine, *para*-benzoylphenylalanine and lysine;
- -X<sub>7</sub> represents an amino acid in the L or D configuration chosen from glycine, alanine, leucine, valine, asparagine and arginine;

 -X<sub>8</sub> represents an amino acid in the L or D configuration chosen from proline, valine, isoleucine and aspartic acid;

 -X<sub>9</sub> represents an amino acid in the L or D configuration chosen from serine, alanine, lysine, arginine and tryptophan;

-the bond between two successive amino acids  $X_i$ - $X_{i+1}$ , denoted  $q_{i \text{ to } i+1}$ , i=1 to 8 can be a peptide

O bond -C-NH - or a pseudopeptide bond chosen from: CO-O, CO-S, CO-CH<sub>2</sub>, CO-N(Me), NH-CO, CH=CH, CH<sub>2</sub>-CH<sub>2</sub>, CH<sub>2</sub>-S, CH<sub>2</sub>-O, CS-NH, CH<sub>2</sub>-NH, CO-CH<sub>2</sub>-NH, CO-NH-NH, CO-NH-N= and CO-N(NH<sub>2</sub>);

-the amino acids stated above X<sub>i</sub>, i = 1 to 9 being capable of comprising a modification of their α-carbon, denoted C<sub>i</sub>, i = 1 to 9 and bearing the side chain R of the amino acid, which modification consisting of the replacement of:

with a group chosen from:

the groups R and CH-R<sub>1</sub> representing the side chain of the amino acid and R<sub>2</sub> representing a  $C_1$ - $C_6$  alkyl group; R-R<sub>2</sub> can constitute a ring,

-the pseudopeptides of the invention also corresponding to the following conditions:

or

one of the bonds  $q_{i \text{ to } i+1}$ , i = 1 to 8 is a pseudopeptide bond

or

one of the C<sub>i</sub>, i = 1 to 9 comprises one of the modifications stated above, wherein said molecule of formula (I) is capable of modulating the proteasome.

Claim 20 (previously presented): The molecule as claimed in claim 19, wherein one or more of the following conditions is verified:

at least one of the integers x0, x1, x2, x4, x7, x8 and x9 is equal to 1;

X<sub>1</sub> and X<sub>3</sub>, which may be identical or different, are chosen from threonine and serine;

X2 is chosen from valine, leucine and isoleucine; or

X<sub>4</sub> is chosen from phenylalanine, tryptophan, tyrosine and para-benzoylphenylalanine.

Claim 21 (previously presented): The molecule as claimed in claim 20, comprising 4 to 8 amino acids.

Claim 22 (previously presented): A molecule as claimed in claims 19 to 21, wherein  $_{x0}$  = 1.

## Claim 23 (canceled)

Claim 24 (previously presented): The molecule as claimed in claim 19, wherein one or more of the following conditions are verified:

- -at least one of X1 and of X2 represents threonine.
- -X2 is chosen from isoleucine and valine,
- -X4 is chosen from phenylalanine, tyrosine and para-benzoylphenylalanine, or
- -at least 2 of the integers x0, x1, x2, x4, x7, x8 and x9 are equal to 1.

Claim 25 (previously presented): The molecule as claimed in claim 19, wherein the molecule corresponds to formula (Ia):

$$X_0$$
- $X_1$ - $X_2$ - $X_3$ - $X_4$ - $X_5$ - $X_6$ 
(Ia)

in which the bonds  $q_{i \text{ to } i+1}$  between the amino acids  $X_i$  and  $X_{i+1}$ , i=1 to 5 are peptide or pseudopeptide bonds.

Claim 26 (canceled)

Claim 27 (canceled)

Claim 28 (currently amended): A molecule of general The molecule as claimed in claim

19, wherein the molecule corresponds to formula (Ib):

in which:

- X<sub>2</sub> represents a natural or synthetic amino acid in the L or D configuration, comprising at least one hydroxyl function on its side chain;
- X<sub>5</sub> represents an amino acid in the L or D configuration chosen from lysine, arginine, histidine, aspartic acid, asparagine, glutamic acid and glutamine;
- X<sub>6</sub> represents an amino acid in the L or D configuration chosen from tyrosine, phenylalanine, leucine, isoleucine, alanine, para-benzoylphenylalanine and lysine;
- X<sub>7</sub> represents an amino acid in the L or D configuration chosen from glycine, alanine, leucine, yaline, asparagine and arginine;
- X<sub>8</sub> represents an amino acid in the L or D configuration chosen from proline, valine, isoleucine
  and aspartic acid;
- X<sub>0</sub> represents an amino acid in the L or D configuration chosen from serine, alanine, lysine, arginine and tryptophan;
- -at least one of the bonds between two successive amino acids is a pseudopeptide bond,

-one of the  $\alpha$ -carbons of one of the amino acids is a modified  $\alpha$ -carbon

Claim 29 (previously presented): The molecule as claimed in claim 19, wherein the molecule is:

CH<sub>3</sub>-(C<sub>n</sub>H<sub>2n</sub>)-CO-TVTYDY with n=4, 6, 8, 10, 12, 14, 16, 18 (SEQ ID NO: 1); CH<sub>3</sub>-(C<sub>n</sub>H<sub>2n</sub>)-CO-TISYDY with n=4, 6, 8, 10, 12, 14, 16, 18 (SEQ ID NO: 2);

or

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CH<sub>3</sub>-(C<sub>n</sub>H<sub>2n</sub>)-CO-TVSYKF with n=4, 6, 8, 10, 12, 14, 16, 18 (SEQ ID NO: 3);
CH<sub>3</sub>-(C<sub>n</sub>H<sub>2n</sub>)-CO-TITFDY with n=4, 6, 8, 10, 12, 14, 16, 18 (SEO ID NO: 4):
CH<sub>3</sub>-(C<sub>n</sub>H<sub>2n</sub>)-CO-TITYKF with n=4, 6, 8, 10, 12, 14, 16, 18 (SEQ ID NO: 5);
CH3-(CnH2n)-CO-TITYEY with n=4, 6, 8, 10, 12, 14, 16, 18 (SEO ID NO: 6);
CH<sub>3</sub>-(C<sub>n</sub>H<sub>2n</sub>)-CO-TITYDF with n=4, 6, 8, 10, 12, 14, 16, 18 (SEQ ID NO: 7);
CH<sub>3</sub>-(C<sub>p</sub>H<sub>2n</sub>)-CO-TVTYKL with n=4, 6, 8, 10, 12, 14, 16, 18 (SEQ ID NO: 8);
CH_3-(C_nH_{2n})-CO-TVTYKY with n=4, 6, 8, 10, 12, 14, 16, 18 (SEQ ID NO: 9);
CH3-(CnH2n)-CO-TVTFKF with n=4, 6, 8, 10, 12, 14, 16, 18 (SEQ ID NO: 10);
CH<sub>3</sub>-(C<sub>n</sub>H<sub>2n</sub>)-CO-TITYDL with n=4, 6, 8, 10, 12, 14, 16, 18 (SEO ID NO: 11):
CH<sub>3</sub>-(C<sub>n</sub>H<sub>2n</sub>)-CO-TVTFDY with n=4, 6, 8, 10, 12, 14, 16, 18 (SEO ID NO: 12);
CH<sub>3</sub>-(C<sub>n</sub>H<sub>2n</sub>)-CO-TVTFKF with n=4, 6, 8, 10, 12, 14, 16, 18 (SEQ ID NO: 13);
CH<sub>3</sub>-(C<sub>n</sub>H<sub>2n</sub>)-CO-TVTYKF with n=4, 6, 8, 10, 12, 14, 16, 18 (SEQ ID NO: 43);
TNL*GPS:
SEK*RVW:
TRA*LVR:
SNL*NDA: or
THI*VIK:
wherein * represents:
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-a bond chosen from ester, thioester, keto methylene, keto methyleneamino, N-methylamide, inverse amide, Z/E vinylene, ethylene, methylenethio, methyleneoxy, thioamide, methyleneamino, hydrazino, carbonylhydrazone and N-amino bonds, or -the presence of an aza-amino acid as a substitution for one of the amino acids adjacent to \*.

Claim 30 (previously presented): The molecule as claimed in claim 19 coupled on its Cterminal end and/or on its N-terminal end with another molecule which promotes its bioavailability.

Claim 31 (previously presented): A composition comprising the molecule as claimed in claim 19 in a pharmaceutically acceptable carrier.

Claim 32 (previously presented): A method for treatment of a disorder or a pathology

associated with proteasome activity comprising administering to an animal in need thereof a molecule as claimed in claim 19.

Claim 33 (previously presented): The method as claimed in claim 32, wherein the disorder or pathology is selected from: cancers involving hematological tumors or solid tumors; autoimmune diseases; AIDS; inflammatory diseases; allograft rejection; and amyotrophy.

Claim 34 (previously presented): A method for radiosensitizing a tumor comprising contacting the tumor with a compound as claimed in claim 19.

Claim 35 (previously presented): A cosmetic and/or dermatological composition comprising a molecule as claimed in claim 19, in a cosmetically and/or dermatologically acceptable carrier.

Claim 36 (previously presented): A cosmetic process for preventing or treating the appearance of effects of chronological skin aging and/or of photoaging, comprising applying to skin the molecule as claimed in claim 19 in a cosmetically acceptable carrier.

Claim 37 (previously presented): The molecule as claimed in claim 21, wherein the molecule comprises 5 to 7 amino acids.

Claim 38 (previously presented): The molecule as claimed in claim 21, wherein the molecule comprises 6 amino acids.

Claim 39 (previously presented): The molecule as claimed in claim 24, wherein at least 3 of the integers  $_{x0}$ ,  $_{x1}$ ,  $_{x2}$ ,  $_{x4}$ ,  $_{x7}$ ,  $_{x8}$  and  $_{x9}$  are equal to 1.

Claim 40 (previously presented): The molecule as claimed in claim 19, wherein p ranges from 2 to 6

Claim 41 (previously presented): The molecule as claimed in claim 19, wherein p ranges

from 5 to 19.

Claim 42 (previously presented): The method as claimed in claim 32, wherein the animal is a human.

Claim 43 (previously presented): The method as claimed in claim 32, wherein the disorder is Alzheimer's disease and Parkinson's disease.

Claim 44 (previously presented): A method for modulating the proteasome of a cell comprising administering the molecule of claim 19 to a cell.

Claim 45 (previously presented): The molecule as claimed in claim 19, wherein  $X_1$  and  $X_3$  both represent threonine.